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(19) (CA) **APPLICATION FOR CANADIAN PATENT** (12)

(54) Device and Method for Separation of Plasma from Blood
and Determination of Blood Analytes

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ABSTRACT OF THE DISCLOSURE

The present invention involves a method and device for obtaining plasma from whole blood and determining one or more analytes contained therein. A sample of blood is first applied to a physical transport medium which moves at least a portion of the blood from a remote location thereon to a second location thereon to a first surface of a microporous plasma separation membrane. This movement is achieved by gravity, absorption or capillary action and the physical transport medium may be a sheet of fabric with weave or otherwise or a channeled or grooved synthetic material. Once the blood has moved to the first surface of the microporous plasma separation membrane, the plasma is absorbed and passes through the membrane, thereby separated from the whole blood (and the red color). The plasma reaches the second (opposite) surface of the microporous plasma separation membrane and contacts a plasma collecting test membrane. This test membrane is capable of reacting with the plasma (i.e. one of its components) to display at least one analyte characteristic thereof. Preferably, colorimetric reaction takes place and color intensity indicates concentration level of the analyte in the blood.

DEVICE AND METHOD FOR SEPARATION
OF PLASMA FROM BLOOD AND
DETERMINATION OF BLOOD ANALYTES
Attorney Docket No. KTI-102A

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BACKGROUND OF THE INVENTION

1. Field of the Invention

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The present invention is directed to a device and a method for separating plasma from blood and determining one or more analytes. More specifically, the present invention is directed to plasma separation from blood and analytical testing of the plasma to measure analytes such as cholesterol, HDL cholesterol, triglycerides, etc.

2. Prior Art Statement

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In many applications such as therapeutics, plasma/serum standards, and particularly for blood analyte determination, there is often a need to separate cells from whole blood. For rapid determination of analytes in ser-

um or plasma, colorimetric methods are quite popular wherein the analyte or a product obtained by conversion of this analyte, reacts with a chromogen to give color, the intensity of which is proportional to the concentration of the analyte. When using whole blood for determination of an analyte, it is essential to remove the red cells from the blood so that the color of the reaction is not masked by the color of the blood. Centrifugation of the blood or the clotting of blood are two common prior art methods used to isolate plasma or serum from whole blood. However, these result in an additional sample preparation step. It would appear that since red blood cells are rather large (about 8 microns in diameter), one could conceivably use a microfilter of a smaller pore size to obtain clear plasma. However, this does not usually work for the following reasons:

(a) When one puts a drop of blood on a dry hydrophilic membrane with pore sizes in the 3 to 8 micron range, the blood gets absorbed into the pores and onto the microfibrils of the filter. Often due to this rapid absorption, hemolysis of blood occurs wherein the fragile red blood cells rupture and the broken cells and/or hemoglobin from the cells leak through. As a result, generally, the plasma coming through is not clear but reddish and the throughput may also be small because the filter may clog up rapidly due to the high solids content of the blood (typically 50% of blood volume is cells). To compound the problem, blood may start clotting in the membrane further promoting the clogging. Due to this clogging, very often large analytes such as lipoproteins may be held back and whatever passes through the membrane may be significantly diluted with respect to these large

analytes. The concentration of these analytes in the filtrate may therefore be significantly lower than in plasma or serum and may vary with both applied sample volume as well as with blood hematocrit (solid content), hence yielding unreliable results.

(b) When one places a drop of blood onto a hydrophilic microporous membrane with pore sizes of less than 3 microns, the clogging on top of the membrane appears to be especially rapid and usually negligible plasma gets through. Again the same problems as mentioned above with clogging are encountered.

(c) Using a depth filter in addition to or in place of the microfilter may likewise be problematic. Due to higher hold up and absorptive capacity of the depth filter, volume requirements of blood become rather excessive and the problems of hemolysis and clogging still

exist. An exception to this is the depth filter made
up of glass fibers which was shown to separate plasma/
serum from blood rather effectively. This method is
described in United States Patent No. 4,477,575. How-
5 ever, many of the problems of other "dry" membrane
systems remain, although to a lesser degree.

In order to overcome the problems of dry membrane
test systems, numerous wet or semi-wet microporous mem-
brane test systems have been proposed. Thus, when
10 wetted with water or with aqueous solutions, the micro-
fibers of these hydrophilic membranes are hydrated by
a thin layer of water. This significantly reduces the
absorptive capacity of the membrane. It is possible to
get fairly clean plasma across this wetted microporous
15 membrane of pore sizes of about 5 microns or less in
this fashion. However, wet and semi-wet systems

have three major drawbacks. First, wet and semi-wet systems may involve wetting the membrane prior to use and this adds an additional

5 step for a diagnostic test. Second, when dealing with small volumes of a few microliters, as are encountered in diagnostic tests, this would dilute the plasma and the extent of dilution would depend upon the volume of sample. Finally, in a wet membrane, the rate of transport
10 of plasma through the membrane will be governed by diffusion rather than capillary pull and may take an excessively long time thereby affecting the overall response time of a diagnostic test.

This approach of using a wet microporous membrane
15 for separation of plasma from whole blood is indeed being used very successfully in a process called plasma-

phoresis. In this process, the microporous membrane is first wetted with isotonic saline in a step which is called priming the membrane. Blood is then pumped through one side of the membrane and a transmembrane pressure differential is maintained. Because of the externally applied pressure difference, the process is fairly rapid. Additionally, a relatively large amount of blood is processed (on the order of a liter) and hence dilution due to isotonic saline in the membrane is negligible. Therefore, even though plasmaphoresis is an effective way of obtaining plasma from blood on a process scale, it is not very practical for a simple and rapid diagnostic test requiring on-line serum/plasma separation for microliter quantities, without external pressures and preferably without dilution.

Glucose reagent strips were among the first to

utilize whole blood for blood glucose measurements.

These colorimetric tests for whole blood based on dry-strip technology became quite popular due to the possibility to obtain rapid analysis of glucose from whole

5 blood without sample pre-treatment. These early tests relied on diffusion of glucose from whole blood through thin, transparent and tightly structured membranes (U.S.

Pats. 3,092,465; 3,298,789; 4,543,338). A drop of blood is placed on the test-pad and after a pre-determined

10 period of time the blood is either washed off or wiped off and the color developed underneath is then compared to a

color chart or read in a reflectance meter. These tests do not require a precise metering of the blood but do require a

blood wash-off or a wipe-off step and precise timing for

15 the same. While this approach is satisfactory for a rela-

tively small hydrophilic molecule such as glucose which can

diffuse rather easily through a thin tightly structured membrane, it is not feasible to measure large analytes such as proteins or hydrophobic analytes such as cholesterol or triglycerides which are complexed with proteins and are carried in the blood stream as part of lipoproteins ranging in size from a few hundred thousand to several million in molecular weights. These relatively large molecules cannot penetrate the dense membranes used in the glucose strips and attempts to separate plasma or serum from whole blood with more open membranes and without pressure or external influences were not successful due to reasons mentioned above.

The discovery that certain depth filters made up of glass fibers could successfully separate red cells from whole blood (U.S. Patent No. 4,477,575) paved the way for successful commercialization of an instrument

called "Reflotron" by Boehringer Mannheim Diagnostics
(Trademark) where the separation of clear serum or
plasma occurs within the dry strip itself without need of
an external force such as pressure or centrifugation.

5 This made analysis of various analytes possible from
whole blood in a simple inexpensive manner. However, al-
though blood analysis with Reflotron is convenient, a metered
amount of blood must be placed onto the reagent pads
and there is a need to crush the pad to bring all the
10 reagents into intimate contact with the separated serum/
plasma. Additionally, it can perform only one reaction
at a time and typically the chemistries take about
three minutes each.

Thus, while many new developments have occurred and
15 have been patented, the need for a convenient, effi-
cient, easy-to-use, reliable test method and device for

testing one or more analytes from whole blood plasma
has not been satisfied until the development of the
present invention.

SUMMARY OF THE PRESENT INVENTION

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The present invention involves a method and device
for obtaining plasma from whole blood and determining
one or more analytes contained therein. A sample of
blood is first applied to a physical transport medium,
which moves at least a portion of the blood from a
remote location thereon to a second location thereon
to a first surface of a microporous plasma separation
membrane. This tangential movement is achieved by gravity,
absorption and/or capillary action. The physical
transport medium may be a sheet of fabric with weave
or otherwise or a channeled or grooved synthetic
material. Once the blood has moved to the first

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surface of the microporous plasma separation membrane,
the plasma is absorbed and passes through the membrane,
thereby separated from the whole blood (and the red
color). The plasma reaches the second (opposite)
5 surface of the microporous plasma separation membrane
and contacts a plasma collecting test membrane. This
test membrane is capable of reacting with the plasma
(i.e. one of its components) to display at least one
--analyte characteristic thereof. Preferably, colori-
10 metric reaction takes place and color intensity
indicates concentration level of the analyte in the
blood.

BRIEF DESCRIPTION OF THE DRAWINGS

The present invention described herein is more
15 fully appreciated and understood when the disclosure
is taken in conjunction with the appended drawings,

wherein:

Figure 1 illustrates a photographic representation of a 5 micron nitrocellulose plasma separation membrane used in the method and device of the present invention;

5 Figure 2 shows a photographic representation of a 3 micron cellulose plasma separation membrane used in the present invention;

10 Figure 3 shows a side cut view of a present invention device which includes a support substrate, a lateral physical transport medium, a microporous plasma separation membrane and a plasma collecting test membrane;

Figure 4 shows a top diagramatic view of the device shown in Figure 3;

15 Figure 5 shows an alternative embodiment present invention device utilizing a channeled substrate as

the physical transport medium;

Figure 6 shows another alternative embodiment
present invention device wherein the physical trans-
port medium is a combined channeled substrate and
capillary action transport membrane;

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Figure 7 illustrates the top view of an altern-
ative present invention device;

Figure 8 illustrates a side view of a present
invention device having multiple plasma collecting
test membranes for determining three different
analytes; and,

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Figure 9 illustrates a present invention device
which includes radially arranged plasma collecting
test membranes which permit analyte determination for
four different analytes.

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DETAILED DESCRIPTION OF THE PRESENT INVENTION

The present invention is directed to obtaining clear plasma from microliter quantities of whole blood rapidly, efficiently, without causing dilution of blood analytes and without relying on externally applied forces or without any sample pretreatment. It is also directed to the process of separation of plasma in a device for determination of blood analytes. The present invention plasma separation method can be utilized for colorimetric detection of blood analytes. The devices of this invention enable the plasma to come into intimate contact with all the necessary enzymes, proteins or other reagents and chromogens without external pressure or without denaturing blood or sensitive reagents. The test strip device of this invention also enables one to do multiple chemistries

on the same strip with a single sample application in
a very quick, simple and reliable manner.

Primarily, the present invention relies upon
application of blood at a first (remote) location on
5 a physical transport medium and movement of the blood
to a second location on the medium for separation of
plasma and subsequent testing. Typically, the present
invention utilizes tangential flow of blood underneath
a suitable microporous plasma separation membrane so
10 that the red cells are retained on the underside but
clean plasma is obtained on the top surface of the
membrane. Capillary pressure, both within the membrane
matrix and by suitable channels and/or a suitable
open mesh underneath the microporous membrane may
15 provide the appropriate driving force for the
tangential flow and no external forces are necessary.

For filtration of highly viscous liquids or for slurries or suspensions with high solids content, tangential flow across the membrane is much more efficient than the dead-ended filtration. The problems of clogging of the membrane are substantially reduced in tangential filtration mode. In this system, a viscous fluid or suspension is swept across one side of the membrane, and clean filtrate is collected across the other side.

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This invention uses this principle, for a diagnostic strip, on a miniature scale. For small volumes of blood which can be obtained by finger-pricking, the driving force for blood flow may be provided entirely by the capillary pull. To do this efficiently, it relies heavily on both the pore-structure of the separation membrane and the physical

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transport medium and support structure underneath it.

Additionally, the choice of the microporous separation

membrane becomes even more important than for other

conventional applications. An ideal separation

5 membrane should wet easily with serum or plasma,

should not allow the red cells to migrate to the top

surface, should not cause hemolysis of the red cells,

should have a good capillary pull and a fairly uniform

pore-structure near the surfaces. Out of several

10 dozen commercial microporous membranes which were

screened, three were found to be particularly preferable,

a 5 micron microporous membrane made from nitrocellu-

lose of Schleicher and Schuell (Keene, New Hampshire)

performed optimally in separating plasma from whole

15 blood. In general, suitable pore-sizes of such mem-

branes should be between about 0.02 to 10 microns, with

the preferred range between 1 to 5 microns. Skinned membranes,

wherein one or both surfaces of the microporous membranes have much smaller and uniform pore-sizes, as compared to the pore-sizes in the middle region, are found to be particularly suitable. Unlike most commercial microporous membranes which are generally isotropic in their cross-section, these membranes show double-skinned structures which may be the reason for their effectiveness. Figures 1 and 2 show the cross-section of the 5 micron nitrocellulose and the 3 micron cellulosic membrane respectively.

Other important components of this invention are the physical transport medium and the substrate onto which this microporous separation membrane is placed. Typically, this membrane is placed on a plastic substrate and between this substrate and the membrane is placed a physical transport medium which may be a material with open structure and hydrophilic surface to achieve tangential flow of the blood.

Examples of materials with such open structures are various types of polymeric meshes, cloth, tissue papers, gauze etc. Ideally the hold up volume and the absorptive capacity of these open-structured materials should be very small so as to minimize the loss of blood due to absorption and the surface on a macroscopic scale should be fairly smooth so as to provide good contact with the microporous separation membrane.

Woven fabrics from many monofilament yarns such as polyester and nylon are particularly useful. Even though the pore sizes of the transport media are not critical, it would be preferable to have them at least an order of magnitude larger than that of the analyte of interest, so that these materials do not have any sieving properties and so that their sole function is to assist in smooth

and rapid flow of blood. As a further aid for rapid movement of blood underneath the microporous separation membrane, channels of well-defined geometry or grooves may be included on the plastic substrate, underneath the open-structured material. Channels 1 to 10 mm wide and 1 to 15 mil (25 to 225 micron) deep work quite well, 1 to 4 mm width and 2 to 10 mil depth being the preferred range.

In practice, an open structured material such as a monofilament woven fabric is placed onto the plastic substrate with or without the channels, and the microporous separation membrane is placed on the top such that part of the fabric extends beyond the membrane. A plasma collecting test membrane such as a piece of filter paper or another hydrophilic membrane is placed on top of the microporous separation membrane. This

will absorb the clean plasma as it comes through the microporous separation membrane and may be loaded with reagents specific for a given analyte. This whole assembly may then be secured by placing another plastic cover on the top. Blood applied to the physical transport medium through an orifice in the plastic cover at a location remote from the separation membrane, migrates along to a location next to (under) the microporous plasma separation membrane, thereby creating a thin film of blood along one surface of the microporous plasma separation membrane. Clean plasma available from the top surface of this membrane saturates the test membrane and if it is loaded with reagents, the analyte reacts with these reagents producing color or other analyte indicator.

Referring now to Figures 3 and 4, there is shown

a side cut view and a top view of present invention
test strip device 1. The device 1 has an inert
substrate 3 and a physical transport medium 5 attached
thereto. In this embodiment, physical transport
5 medium 5 is a woven cloth material. Atop physical
transport medium 5 is microporous plasma separation
membrane 7 and optional hydrophobic barrier strip 9,
adjacent thereto as shown in both figures. Of less
length and placed atop of microporous membrane 7 is
10 plasma collecting test membrane 11. Clear plastic
cover 13 with orifice 15 is adhered over the tops of
the other components as shown by the arrows 2 and 4
in Figure 3. Top view Figure 4 shows all of the
components, at least in part, due to the fact that
15 cover 13 is clear.

As now can be seen, the user simply places a drop

of blood in orifice 15 (a remote location from
microporous plasma separation membrane 7). The
blood flows down physical transport medium 5 and under
microporous plasma separation membrane 7 to a first
5 (bottom) surface thereof. Plasma separates and flows
upwardly through membrane 7 and is, at this point,
essentially clear in color. It migrates to a second
(opposite, i.e. top) surface of membrane 7 to plasma
collecting test membrane 11. Here the analyte(s) react
10 to establish concentration(s). For example, in this
case , test membrane 11 contains reagents specific
for the analyte or analytes of interest, e.g.
chromogens which reveal colorimetric measurement of
cholesterol, glucose or the like. Further, the
15 dimensions shown in Figure 4 are merely exemplary and
are not critical.

In general, the essential components of the present invention device are the physical transport medium, the microporous plasma separation membrane, and the plasma collecting test strip. Although not preferred, a microporous plasma separation membrane may be extended and graded to act both as a transport medium and a separation membrane. However, more leeway as to blood volume, dimensions and "overflow" control are achieved when the physical transport medium is one or more components which are elements separate and different from the microporous plasma separation membrane. Likewise, the transport medium may act as a substrate or a separate, inert and inactive substrate may be utilized.

The combination of a separate physical transport medium and a separation membrane is chosen such that

when blood is placed at the remote blood application area, it is rapidly pulled underneath the microporous membrane and clean plasma seeps across this membrane and saturates the reactant containing test membrane.

5 Further, the test membrane may be a single layer or may be a compilation e.g., a reagent containing layer as well as a chromogen containing layer which is preferably a porous hydrophilic layer. The enzymes or
10 ~~reagents in these layer(s)~~ dissolve in the plasma and react with the analyte producing color. This type of system is particularly appropriate for determining analytes such as glucose, cholesterol, triglycerides, uric acid, creatinine, alcohol, creatine kinase, cholinesterase, AST, ALT, etc. wherein an enzyme or
15 a series of enzymes convert an analyte to hydrogen peroxide which then reacts with chromogen, contained

in the test membrane. Many of the enzymes used in
such diagnostic strips are particularly susceptible to
denaturation when contacted by many chromogens for
an extended period of time. By separating the enzymes
5 and the chromogens in two separate layers and yet
keeping them in close proximity, the long term
stability of the reagents can be preserved. For
analyzing analytes such as albumin or other proteins,
amylases, calcium, inorganic phosphorus, etc. wherein
10 the reagents used are not biomolecules such as enzymes
or antibodies but are stable chemicals and the color
formed is due to selective complexation of the analyte
with the reagents, the porous layer can be eliminated
and all the reagents can be combined in a single layer
15 membrane. It is also possible to load some of the
reagents elsewhere in the system such as in additional

layers, although this is not essential, provided that they don't hemolyze the cells or damage them in any way.

It is best to view the final color at the top of the test membrane since it also acts as a hiding layer to

5 mask the color of the blood. In such a case, an additional layer, if necessary for chromogens, could be transparent or translucent. In order to avoid the bleedthrough of the color developed during the test

10 ~~downstream from the device onto the test membrane, an~~ optional hydrophobic material or glue can be applied on the exposed edge of the separation membrane just downstream of the physical transport medium (not shown in the Figures). Such a hydrophobic surface barrier prevents the color from diffusing down the separation
15 membrane, but is not always necessary.

In the alternative, when a test membrane

having a first reactant (reagent) and second reacting component (chromogen) is used, a single test membrane instead of a plural layer may be used. Thus, in some embodiments, the reagents and chromogens may be contained in dry or semi-wet form or even wet form in the same membrane structure, or one or the other may be encapsulated or otherwise isolated within the same membrane.

Further, the present invention device may be used without any additional sample preparation for detection of certain analytes which normally requires a precipitation step for removal of interfering analytes. The precipitants, for example, can be incorporated within the microporous plasma separation membrane or in the physical transport medium, so that the precipitation step occurs on-line and the plasma reaching the test

membrane is free of the interferent. If, due to the nature of the precipitant, it is advisable not to have direct contact of the precipitant with the blood, then an additional layer containing the precipitant can be placed between the separation membrane and the physical transport medium. An application that makes use of the precipitation step is the measurement of high density lipoprotein cholesterol (HDL cholesterol) which typically needs precipitation steps using precipitants such as polyethylene glycols, dextran sulfate, or magnesium tungstate to precipitate out cholesterol in low density and very low density lipoproteins (LDL and VLDL), followed by centrifugation. The present invention device achieves this on-line by loading the precipitant into the microporous plasma separation membrane.

These devices are self-monitoring because a microporous membrane cannot be over-filled or over-saturated by capillary filling. Once the plasma collection membrane is saturated, the excess blood and plasma simply migrate down the transport medium. If, for some reason, the sample volume is so large as to exhaust the absorptive capacity of the transport medium, then the excess blood will merely stay at the application area without getting absorbed. In case the sample volume is not enough, then only part of the test membrane will be saturated and colored. The precise control over the thicknesses (and hence the void volume) of the test membrane is not critical because even if the saturation capacity of this membrane varies with the thickness, the concentration of the analyte and hence the intensity of the final

color produced will be independent of the thickness of this membrane. The minimum volume needed to saturate the plasma collecting test membrane with plasma will depend on the thickness of this membrane, the geometry of the system and the hematocrit of the blood: for the system shown in Figures 3 and 4, it is approximately 10-15 microliters. When using plasma or serum in place of whole blood, the volume requirements are half as much. To further minimize the required minimum volume the size of the test membrane and/or the thickness may be decreased as long as the test membrane is sufficiently opaque to mask the color of the blood underneath it. If the test membrane is not of sufficient opacity, then one could use a geometry similar to that shown in Figure 7 where the color developed is not directly over the blood path.

Figure 5 shows a present invention device 21,
which is similar to the one shown in Figures 3 and 4
and like parts are like numbered. Note that physical
transport medium 19 is now channels or grooves
5 established in substrate 17 itself. Arrows 6 and 8
show assemblage.

Figure 6 likewise shows a device 23 with like
parts being identically numbered, and here, both the
cloth physical transport medium 5 of the Figure 3
10 device and the grooved physical transport medium 19
of the Figure 5 device are combined. Additionally,
absorbant storage medium 25 rests in part on transport
medium 5 and on substrate 17. This optional storage
medium may be used to absorb additional (excess)
15 blood to draw the excess past the separation membrane
7. Any blood absorbing material may be used, although

this is not an essential feature of the invention.

Figure 7 shows present invention device 33 which has substrate 35, physical transport medium 37 (e.g. a sheet material such as a gauze or synthetic having the necessary capillary action), located on substrate 35, plasma separation membrane 39 over physical transport medium 37, and plasma collecting test membrane 41 which is partly over the separation membrane 39 and partly located directly on substrate 35, as shown. This arrangement permits the use of a very thin separation membrane which may show through some of the red color. By offsetting part of the test membrane, the isolated portions will show true colorimetric readings without a pinkish background. Top cover 43 with orifice 45 is clear plastic and the user uses this strip in the same manner as described

above.

Advantageously, the present invention method and device may be used for a plurality of simultaneous testings. Thus, it is possible to combine various chemistries on the same device. This is particularly advantageous for example in testing on-site in doctor's office, where a patient needs to be checked for several analytes to diagnose a condition. In the strips of this invention several related tests can be combined on the same strip and determination of multiple analytes can be made within a few minutes from application of a single blood sample which can be obtained by finger-prick. Thus, Figure 8 shows one geometry combining a lipid panel (cholesterol; HDL cholesterol and triglycerides) onto a single strip. There is shown in Figure 8 device 41 having substrate

43. The physical transport medium in this embodiment includes channels 45 located on substrate 43 as well as sheet 47 located above channels 45. Three microporous plasma separation membranes 49, 51 and 53 are spaced apart and located on sheet 47 as shown. Membrane 51 additionally contains a precipitant for LDLs and VLDLs (low and very low density cholesterol). Atop each of the separation membranes 49, 51 and 53 are multilayer plasma collecting test membranes 55, 57 and 59 respectively. The plasma collecting test membranes 55, 57 and 59 contain the necessary reactants for three different analytes, in this case cholesterol, HDL cholesterol and triglycerides respectively. Optional absorbant storage medium 61 is also included. The top cover in this embodiment includes a first clear plastic layer 63 with orifice 65 and an opaque layer 67 containing orifice 69 for blood

sample transmission and orifices 71, 73 and 75 for visual comparison for colorimetric measurements of the respective analyte reactions at test membranes 55, 57 and 59.

Figure 9 shows a different geometry for obtaining plural analyte characteristics and, in this case, contains four different analyte tests. Thus, device 91 includes substrate 93 and, in a large radially expanding cross, physical transport medium 95 which is placed on top of substrate 93 and extends radially from the center to approximately the outer edge of substrate 93 and under all other components hereafter described. Atop physical transport medium 95 are four separate microporous plasma separation membranes 97, 99, 101 and 103. Atop a portion of each of these are plasma collecting test membranes 105, 107, 109 and 111, respectively, and these in turn contain reactants for colorimetric determination for four different analytes.

such as glucose, cholesterol, HDL cholesterol and triglycerides. Optional absorbant storage media 113, 115, 117 and 119 are placed on top of the outer ends of physical transport medium 95. Clear plastic cover 121 is placed on top of all the other components and has a single orifice 123 for blood sample application in the center as shown. This device 91 permits nearly simultaneous testing of four different analytes by application of a single blood drop.

10 The following examples further support the effectiveness of the present invention:

Example 1

 A device such as that shown in Figures 3 and 4 was prepared except that tissue paper is used as the
15 the physical transport medium to facilitate the flow of blood underneath a 5 micron pore-size nitro-

cellulose separation membrane (Type AE 98, Schleicher and Schuell), a microphotograph of which is shown as Figure 1. The plasma collecting test membrane in this case was a 0.45 micron nylon 6,6 membrane. It is advantageous to use for the test membrane a microporous membrane of a pore size smaller than the plasma separation membrane for plasma collection; it will then also act as a secondary membrane in case the primary membrane has some defects. A drop of blood (20-40 microliters in volume) is placed at the blood application area and within about 1 to 2 minutes the plasma collection membrane gets saturated with clear plasma.

Example 2

A Figure 5 type of device was constructed wherein the bottom support plastic has a 3 mm wide and 5 mil deep channel built in. No open structured material is

used as a blood transport material. To prevent the blood from going over the plasma separation membrane, a hydrophobic tape is put on the top surface of this membrane. The plasma separation membrane is the very same as in Example 1. When a drop of blood is placed on the blood application area, clear plasma saturates the plasma collecting test membrane in about 1 minute.

Example 3

A Figure 6 device with a geometry utilizing both the channels and an open mesh was made. The channel geometry is the same as in Example 2 but a woven fabric from monofilament polyester is used to facilitate the transport of blood. The plasma separation membrane is of the same type as in above examples. When a drop of blood is placed on the blood application area, clear plasma saturates the plasma collection membrane in

about 10 seconds.

Example 4

The same geometry as shown in Example 3 is used except that the plasma separation membrane is 8 micron nitrocellulose (Type AE99, Schleicher and Shuell). The plasma collecting test membrane is saturated in about 10 seconds.

Example 5

The same geometry as in Example 4 is used except that cellulosic membrane (Micron Separations, Inc.) is used for the separation membrane. A microphotograph of this cellulosic membrane is shown in Figure 2. The plasma collecting membrane is saturated within 30 seconds.

Example 6

In the geometry of Example 3, an additional filter paper is introduced to serve as an absorption pad to soak up excess blood as shown in Figure 7. The rest of

the components are the same as in Example 3. Blood samples from about 15 to 100 microliters are placed in the blood application area and the plasma collection membrane is saturated in about 10 seconds.

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Examples 7-15

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Examples 7 through 15 are illustrative of the geometry, assembly and functioning of the devices. In these illustrative examples, a 5 micron nitrocellulose membrane (Schleicher and Schuell) was used as the plasma separation membrane (#1), 0.45 micron pore-size nylon 6.6 membrane (Micron Separation Inc., Westoro, MA) as the plasma collecting test membrane and a woven polyester mesh as the physical transport medium to facilitate the transport of blood, lens tissue-paper or a dialysis membrane of 10,000 dalton cut-off (Spectrum Medical Industries, Inc., L.A., CA) was used to load the chromagen, white PVC or polystyrene

plastic (15-25 gauge) was used as the substrate and
a clear PVC or vinyl plastic (8-12 gauge) was used as
the top cover. Glues such as polybutene or silastic
or PVC based, or medical grade double-stick tape, all
5 of which are relatively inert toward the chemistries,
were used in these devices to provide either a hydro-
phobic barrier or a good seal. The chromogen used in
conjunction with the various chemistries involving
enzymes as the reagents is 3,3', 5,5' tetramethyl-
10 benzidine dihydrochloride (TMBD) which in the presence
of horseradish peroxidase gives bluish green color
when it reacts with hydrogen peroxide generated from
the reactions of the analyte. TMBD can be substituted
by a number of other chromogens such as phenol or its
15 derivatives and 4 amino antipyrine, or a number of
leuco dyes, e.g. from arylmethine family (e.g. leuco

malachite green) or by solubilized reactive dyes in their leuco form. The porous lower layer of the test membrane was soaked into the solution of the chromogen and allowed to dry. Typically this solution

5 consisted of 0.3g TMBD, 0.5g of sodium cholate, and 0.5-2g of polyethylene glycol (PEG) (molecular weight 8000 daltons) dissolved in 20 ml of reagent alcohol and 0.1M pH 3 phosphate buffer (3:1 by vol). The top layer of the test membrane was tissue paper soaked in

10 the chromogen solution and upon drying stays tacky due to a combination of sodium cholate and PEG. This tackiness enables it to have an intimate contact with the reagent lower layer. To load the reagents, reagent solutions in appropriate buffers are made at

15 the desired concentrations and the membrane is either soaked into this solution or the solutions pipetted

onto the membrane till it is saturated. The membrane is then allowed to dry at room temperature. Typically the reagent membrane needs 10 microliter of reagent solution per square cm area for saturation. The

5 solutions used for the various tests illustrated in the examples below are as follows: (a) cholesterol: 400 units of pancreatic cholesterol esterase, 40 units of cholesterol oxidase from Nocardia, and 200 units of horseradish peroxidase per milliliter of 0.1M
10 pH 7 phosphate buffer. (b) Triglycerides: 2000 units of lipase from chromobacterium viscosum, 300 units of glycerol kinase from arthrobacterium sp., 300 units of glycerol phosphate oxidase (Microbial), and 500 units
15 of horseradish peroxidase per milliliter of 0.1M phosphate buffer at pH 8. Additionally adenosine triphosphate (ATP) solution is prepared at a concentration of .625g/ml and

is combined with the above solution in the ratio of
4 parts of enzyme solution to 1 parts of ATP
solution. (c) glucose: 80 units of glucose oxidase
from aspergillus niger and 300 units of horseradish per-
oxidase per milliliter of 0.1M phosphate buffer at pH 7.

Example 7

Test strips for measurement of total cholesterol
were made as per Figure 6 where the reagent membrane
contained the enzymes for cholesterol as described
above. Blood or serum samples ranging in volume from
10 to 40 microliter were placed at the blood
application area. Typically it took about 10 to 15
seconds to saturate the test membrane and in about
30 seconds the color was fully developed which was
then read in a reflectance meter or compared to a
color chart.

Example 8

Test strips of cholesterol were made as per the geometry of Figure 6 and with the reagent composition as described above. Samples of whole blood ranging in volume from 15 to 50 microliters were applied at the blood application area. The color development typically was complete in about 30 to 45 seconds.

Example 9

HDL cholesterol strips were made as per geometry of Figure 6. The only difference between the cholesterol strips of Example 7 and the HDL strips is that the plasma separation membrane 1 in this case was first soaked in an aqueous 20% by weight solution of PEG 2000 (a precipitating agent for LDL and VLDL cholesterol) and allowed to dry prior to use. 10 to 40 microliter samples (whole blood, serum or plasma)

were applied at the sample application area and color development was complete in about 1 minute.

Example 10

5 Triglyceride strips were made as per the geometry of Figure 6 and the test membrane was loaded with the enzymes as described above. 10 to 40 microliter samples (whole blood, serum or plasma) were applied at the sample application area and color development was complete in about 1 minute.

10

Example 11

Glucose test strips were made as per the geometry of Figure 6 wherein the test membrane was loaded with the enzymes as described above. 10 to 40 microliter samples (whole blood, serum or plasma) were applied at
15 the sample application area and color development was complete in about 30 seconds.

Example 12

Test strips for the lipid measurement were prepared as shown in Figure 8. The three test membranes for individual geometries were as described above and the separation membrane for HDL cholesterol was treated with PEG as in Example 9. 40 to 100 microliter samples (whole blood, serum or plasma) were applied at the sample application area and color development was complete in about 3 minutes.

Example 13

Devices for simultaneous measurement of cholesterol (total and HDL), glucose and triglycerides was prepared as per the geometry of Figure 9. 50 to 100 microliter samples (whole blood, serum or plasma) were applied at the sample

application area and color development was complete
in about 2 minutes.

Example 14

5 Devices for the measurement of total cholesterol
and HDL cholesterol were made. 20 to 80 micro-
liter samples were applied at the sample application
area and the color development was complete in
about 1 minute.

Examples 15-21

10 In the following Examples 15 - 21, the
enzymes and chromogens were loaded in a single
layer plasma collecting test membrane. The
chromogen was 3,3', 5,5' tetramethyl benzidine
which was dissolved at a concentration of .03 g/ml
15 in acetone or at about .02 g/ml in a 3% solution

of phenol in toluene. The enzymes for cholesterol
were loaded from their aqueous solutions as
described above and the test membranes were allowed
to dry. After drying, the test membranes were
5 loaded with chromogen solution (either toluene
or acetone based) and again allowed to dry.
The geometry used in testing was as shown in
Figure 7, except that the separate chromogen
layer was eliminated and additionally an aperture
10 was provided on the top plastic cover over the
plasma collecting membrane to allow for oxygen,
since the reactions seemed more sensitive to
oxygen requirements than when the chromogen
was loaded separately in another matrix. For
15 aesthetic purposes and to protect the
reagent area from outside contamination,

an air-permeable membrane, such as a
thin (~20 micron thick), transparent polyurethane
membrane coated on one side with porous adhesive (e.g.
from Acutek, Inglewood, California) can be placed
5 over the aperture. Such films were found to allow
adequate transport of oxygen to the reaction zone.

Example 15

A nylon -6,6 membrane of 0.45 micron pore-size
was loaded with cholesterol enzymes and chromogen
10 from acetone solution. When blood or serum samples
in volumes ranging from 15 to 50 microliters were
applied at the blood application zone, in a Figure 7
type device, color developed in about 30 seconds and
colors were accurately proportional to the cholesterol
15 content.

Example 16

The same test was done as in Example 16 with a cellulosic test membrane of 0.2 micron pore-size. The chromogen, however, was loaded from its solution in toluene. Again similar results were obtained.

5 Example 17

The test of Example 15 was repeated. This time, however, after loading and drying with cholesterol enzymes, the test membranes were saturated with acetone, and after partial evaporation of acetone, a mixture containing 90:10 parts by volume of toluene and dodecane was added. After a few minutes of drying the test membrane, chromogen in toluene solution was applied. When the test membrane dries, the non-volatile dodecane component is left behind. These membranes are, therefore, not quite dry but contain the hydrophobic, non-polar dodecane and can be

10

15

considered pseudo-liquid membranes. Non-polar liquids
such as dodecane create a hydrophobic environment
around the enzymes and give them added stability
against denaturation and especially under humid
5 environment. When tested, the performance was
similar to above.

Example 18

The test of Example 15 was repeated but this
time with a test membrane from HYPAN[®] multiblock
10 polymer (Kingston Technology, Dayton, N.J.) and
described in U.S. Patents Nos. 4,331,783; 4,337,327;
4,369,294; 4,379,874; 4,420,589 and 4,370,451. This
test membrane was made over a polyester cloth backing
and contained approximately 5-10 mole % of acrylamide
15 groups and the remainder were acrylonitrile groups.
The structure of the test membrane was asymmetric with

the pores on the more open side of being larger than 1 micron. When this membrane was tested as in Example 15, similar results were obtained.

Example 19

5 The same test as that in Example 17 was repeated with HYPAN membrane of Example 18. HYPAN membrane in this pseudo-liquid state showed similar behavior.

Example 20

10 The test of Example 15 was repeated with a microporous HYPAN membrane which was symmetric in structure. The membrane was impregnated on a polyester cloth support. The HYPAN membrane in this case had greater than 75 mole % of acrylonitrile groups, the remainder being glutarimide-derived groups.

15 The pores on this membrane were in the range of .02-1 micron. The test results were similar to those in

Example 15.

Example 21

The membrane of Example 20 was tried in a pseudo-liquid form as described in Example 17. Again
5 similar successful results were obtained.

Obviously, numerous modifications and variations of the present invention are possible in light of the above teachings. It is therefore understood that within the scope of the appended claims, the invention may be practiced otherwise than as specifically
10 described herein.

The embodiments of the invention in which an exclusive property or privilege is claimed, are defined as follows :

1. A method of obtaining plasma from whole blood

and determining blood analytes thereof, which

comprises:

(a) applying a sample of blood, directly or

indirectly, to a first surface of a micro-

porous plasma separation membrane to effect

separation of plasma from the whole blood

and to effect movement of the separated

plasma to a second surface of said micro-

porous plasma separation membrane;

(b) contacting a plasma collecting test membrane

with said separated plasma, said test

membrane being capable of reacting with said

plasma to display at least one analyte

characteristic thereof.

2. The method of claim 1 wherein said sample of blood is indirectly applied to said first surface of said microporous plasma separation membrane by first being applied to a physical transport medium which moves at least a portion of said blood along the physical transport medium from a remote lock on said physical transport medium to said first surface of microporous plasma separation membrane.

3. The method of claim 2 wherein said physical transport medium relies, at least in part, on capillary forces to move the blood tangentially.

4. The method of claim 2 wherein said physical transport medium is a transport membrane sheet.

5. The method of claim 3 wherein said physical transport medium is a transport membrane sheet.

6. The method of claim 4 wherein said transport membrane sheet is a sheet of fabric selected from woven fabric, non-woven fabric, gauze and monofilament yarn.

7. The method of claim 5 wherein said transport membrane sheet is a sheet of fabric selected from woven fabric, non-woven fabric, gauze and monofilament yarn.

8. The method of claim 2 wherein said physical transport medium constitutes grooves or channels formed on an inert substrate to promote movement of blood.

9. The method of claim 3 wherein said physical transport medium constitutes grooves or channels formed on an inert substrate to promote movement of blood.

10. The method of claim 2 wherein said physical transport medium is an open-structured mesh with openings of at least 1 micron in size and having hydrophilic surfaces.
11. The method of claim 3 wherein said physical transport medium is an open-structured mesh with openings of at least 1 micron in size and having hydrophilic surfaces.
12. The method of claim 1 wherein said microporous plasma separation membrane has a nominal pore size of about 0.02 to about 10 microns.
13. The method of claim 2 wherein said microporous plasma separation membrane has a nominal pore size of about 0.02 to about 10 microns.

14. The method of claim 3 wherein said microporous plasma separation membrane has a nominal pore size of about 0.02 to about 10 microns.
15. The method of claim 1 wherein said microporous plasma separation membrane is a skinned membrane.
16. The method of claim 2 wherein said microporous plasma separation membrane is a skinned membrane.
17. The method of claim 3 wherein said microporous plasma separation membrane is a skinned membrane.
18. The method of claim 1 wherein said microporous plasma separation membrane is cellulosic.
19. The method of claim 2 wherein said microporous plasma separation membrane is cellulosic.

20. The method of claim 3 wherein said microporous plasma separation membrane is cellulosic.
21. The method of claim 1 wherein said microporous plasma separation membrane is of nitrocellulose.
22. The method of claim 2 wherein said microporous plasma separation membrane is of nitrocellulose.
23. The method of claim 3 wherein said microporous plasma separation membrane is of nitrocellulose.
24. The method of claim 2 wherein said physical transport medium moves excess blood to an absorbent storage medium.
25. The method of claim 3 wherein said physical transport medium moves excess blood to an

absorbent storage medium.

26. The method of claim 1 wherein said plasma collecting test membrane includes one or more reactants which react with said plasma to effect colorimetric reactions of one or more analytes.
27. The method of claim 2 wherein said plasma collecting test membrane includes one or more reactants which react with said plasma to effect colorimetric reactions of one or more analytes.
28. The method of claim 3 wherein said plasma collecting test membrane includes one or more reactants which react with said plasma to effect colorimetric reactions of one or more analytes.
29. The method of claim 1 wherein said analyte

characteristic is a colorimetric reaction result.

30. The method of claim 2 wherein said analyte characteristic is a colorimetric reaction result.

31. The method of claim 3 wherein said analyte characteristic is a colorimetric reaction result.

32. The method of claim 1 wherein said analyte characteristic is analyte concentration of one or more analytes selected from the group consisting of cholesterol, HDL cholesterol, triglycerides and glucose.

33. The method of claim 2 wherein said analyte characteristic is analyte concentration of one or more analytes selected from the group

consisting of cholesterol, HDL cholesterol, triglycerides and glucose.

34. The method of claim 3 wherein said analyte characteristic is analyte concentration of one or more analytes selected from the group consisting of cholesterol, HDL cholesterol, triglycerides and glucose.
35. A device for obtaining plasma from whole blood and determining blood analytes thereof, which comprises:
- (a) a microporous plasma separation membrane for applying a sample of blood, indirectly, to a first surface of said microporous plasma separation membrane to effect separation of plasma from the whole blood

and to effect movement of the separated plasma to a second surface of said microporous plasma separation membrane;

(b) a plasma collecting test membrane in contact with the second surface of said microporous plasma separation membrane for receiving said separated plasma, said test membrane being capable of reacting with said plasma to display at least one analyte characteristic thereof; and,

(c) a physical transport medium contacted in part with said first surface of said microporous plasma separation membrane which medium is capable of moving at least a portion of blood to be tested

from a remote location on said physical
transport medium to said first surface
of said microporous plasma separation
membrane.

36. The device of claim 35 wherein said physical
transport medium relies, at least in part, on
capillary forces to move the blood.
37. The device of claim 35 wherein said physical
transport medium is a transport membrane sheet.
38. The device of claim 36 wherein said physical
transport medium is a transport membrane sheet.
39. The device of claim 37 wherein said transport
membrane sheet is a sheet of fabric selected
from woven fabric, non-woven fabric, gauze and
monofilament yarn.

40. The device of claim 38 wherein said transport membrane sheet is a sheet of fabric selected from woven fabric, non-woven fabric, gauze and monofilament yarn.
41. The device of claim 35 wherein said physical transport medium constitutes grooves or channels formed on an inert substrate to promote movement of blood.
42. The device of claim 36 wherein said physical transport medium constitutes grooves or channels formed in an inert substrate to promote movement of blood.
43. The device of claim 35 wherein said physical transport medium is an open-structured mesh with openings of at least 1 micron in size and having

hydrophilic surfaces.

44. The device of claim 36 wherein said physical transport medium is an open-structured mesh with openings of at least 1 micron in size and having hydrophilic surfaces.

45. The device of claim 35 wherein said microporous plasma separation membrane has a nominal pore size of about 0.02 to about 10 microns.

46. The device of claim 36 wherein said microporous plasma separation membrane has a nominal pore size of about 0.02 to about 10 microns.

47. The device of claim 37 wherein said microporous plasma separation membrane has a nominal pore size of about 0.02 to about 10 microns.

48. The device of claim 35 wherein said microporous plasma separation membrane is a skinned membrane.

49. The device of claim 35 wherein said microporous plasma separation membrane is a cellulosic.

50. The device of claim 35 wherein said microporous plasma separation membrane is of nitrocellulose.

51. The device of claim 35 further comprising:

- (d) an absorbent storage medium connected to said physical transport medium wherein said physical transport medium moves excess blood to said absorbent storage medium to prevent excess blood from retarding the effectiveness of said plasma collecting test membrane.

52. The device of claim 37 wherein an absorbent storage medium connected to said physical transport medium wherein said physical transport medium moves excess blood to said absorbent storage medium to prevent excess blood from retarding the effectiveness of said plasma collecting test membrane.

53. The device of claim 38 wherein an absorbent storage medium connected to said physical transport medium wherein said physical transport medium moves excess blood to said absorbent storage medium to prevent excess blood from retarding the effectiveness of said plasma collecting test membrane.

54. The device of claim 35 wherein said plasma collecting test membrane includes one or more reactants which react with said plasma to effect colorimetric reactions of one or more analytes.
55. The device of claim 35 further including at least one inert substrate to which at least one of said microporous plasma separation membrane, said plasma collecting test membrane and said physical transport medium is connected.
56. A device for obtaining plasma from whole blood and determining blood analytes thereof, which comprises:
- (a) an inert substrate support;
 - (b) a physical transport medium attached to a surface of said support;

- (c) a microporous plasma separation membrane having a first surface against said physical transport medium and having a second surface opposite and away from said physical transport medium; and,
- (d) at least one plasma collecting test membrane in intimate contact with a portion of said second surface of said microporous plasma separation membrane and having reactants therein capable of reacting with said plasma to display at least one analyte characteristic thereof.

57. The device of claim 56 wherein a portion of said physical transport medium with which the microporous plasma separation membrane is not in contact, is exposed for applying a sample of

blood thereto.

58. The device of claim 56 which further comprises:

(e) a top sheet containing at least one aperture through which a blood sample may be applied onto the physical transport medium, and at least a portion of this cover sheet overlying said plasma collecting test membranes, said portion being transparent areas to view one or more colorimetric reactions.

59. The device of claim 58 wherein portions of said transparent areas on the cover sheet, have holes or apertures to ensure oxygen transport to the plasma collecting test membrane for reactions that utilize oxygen.

60. The device of claim 59 wherein said holes or apertures for oxygen transport are covered by transparent oxygen permeable membranes.



FIG. 1



FIG. 2

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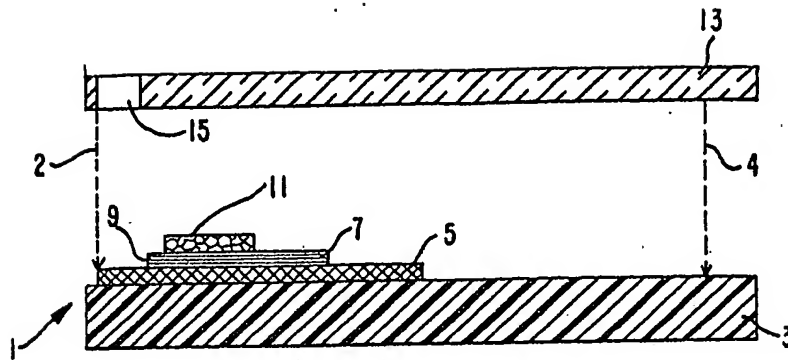


FIGURE 3

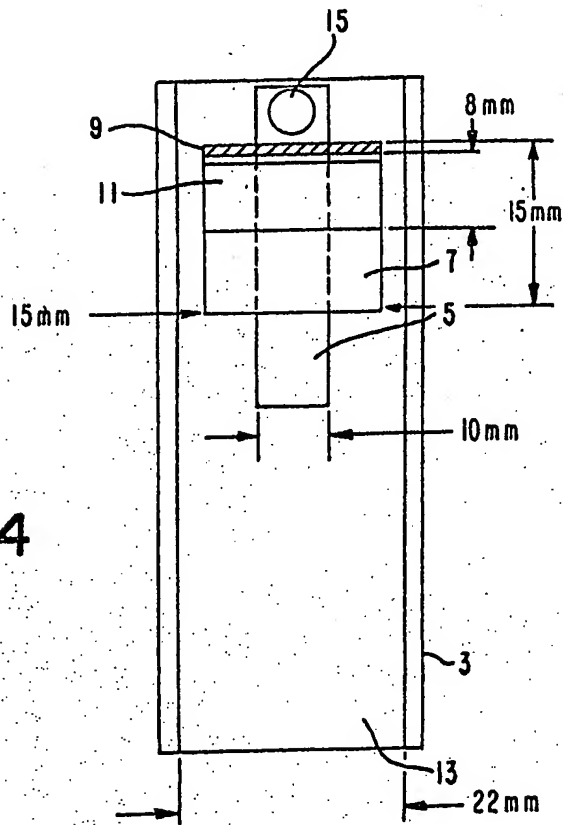
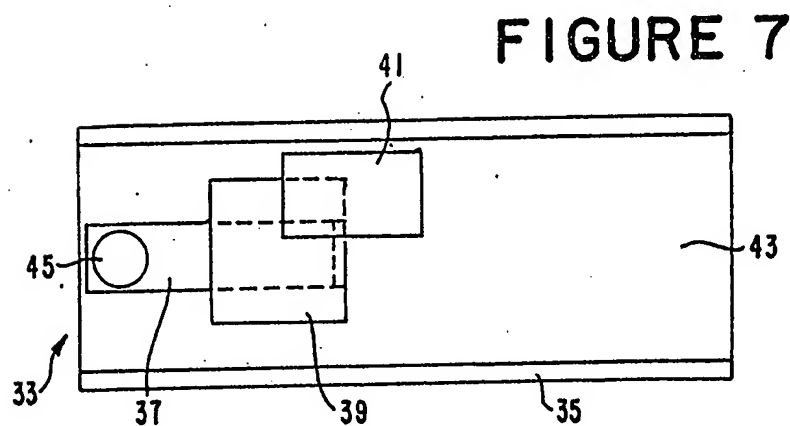
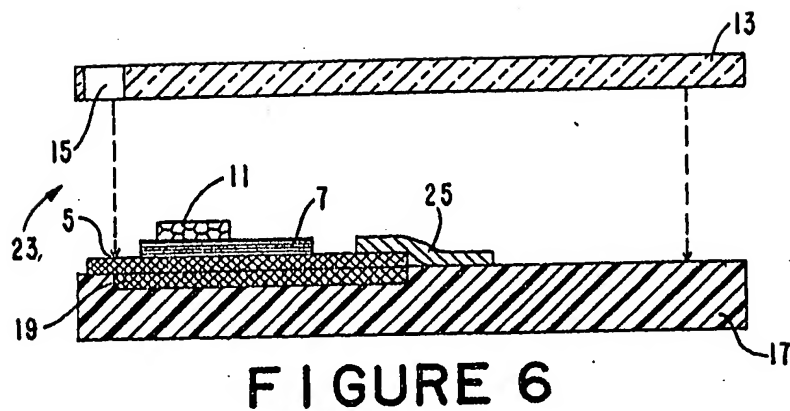
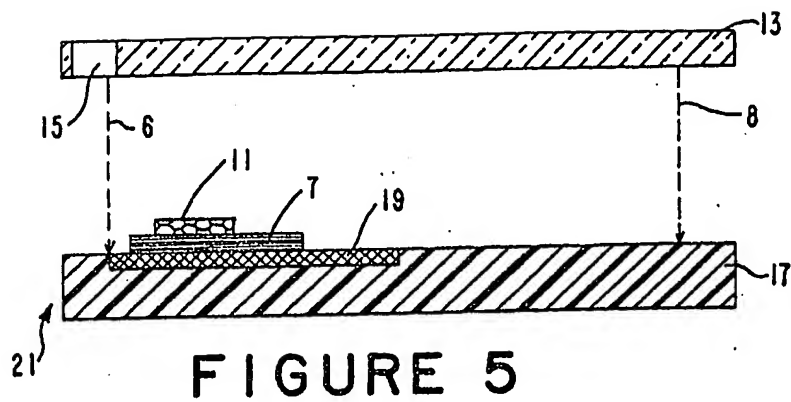


FIGURE 4

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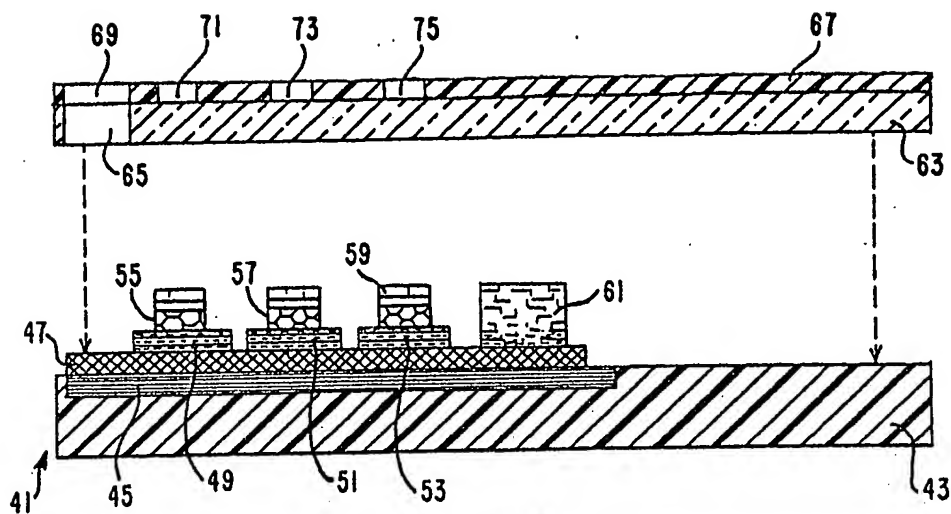


FIGURE 8

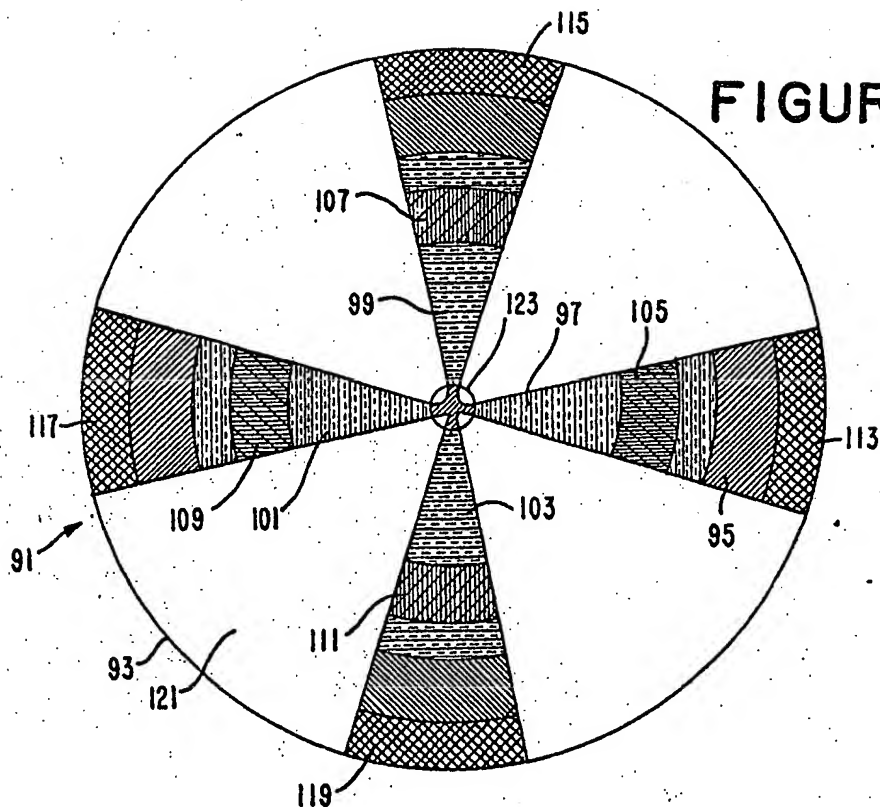


FIGURE 9

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